

---

# DICTIONARY OF BIOCHEMISTRY AND MOLECULAR BIOLOGY

Second Edition

**J. STENESH**

*Professor of Chemistry  
Western Michigan University*



WILEY

A WILEY-INTERSCIENCE PUBLICATION

**JOHN WILEY & SONS**

New York / Chichester / Brisbane / Toronto / Singapore

---

Copyright © 1989 by John Wiley & Sons, Inc.

All rights reserved. Published simultaneously in Canada.

Reproduction or translation of any part of this work beyond that permitted by Section 107 or 108 of the 1976 United States Copyright Act without the permission of the copyright owner is unlawful. Requests for permission or further information should be addressed to the Permissions Department, John Wiley & Sons, Inc.

*Library of Congress Cataloging in Publication Data:*

Stenesh, J., 1927—

Dictionary of biochemistry and molecular biology / J. Stenesh. —  
2nd ed.

p. cm.

Rev. ed. of: Dictionary of biochemistry, 1975.

"A Wiley-Interscience publication."

Bibliography: p.

ISBN 0-471-84089-0

1. Biochemistry—Dictionaries. 2. Molecular biology—  
—Dictionaries. I. Stenesh, J., 1927— Dictionary of biochemistry.

II. Title.

QP512.S73 1989

574.19'2'0321—dc19

88-38561

CIP

Printed in the United States of America

10 9 8 7 6 5 4 3 2

This dictionary  
ten to provide  
sciences with a  
logy of biochem  
expansion of k  
the need for a  
edition. All of  
and reworked,  
formation. Thi  
imately 16,000  
new, represent  
over that of th  
rial consulted f  
for addition of  
textbooks and  
and of over 6  
search literatur  
lished since 197  
are drawn from  
icles, including  
Commission of  
the International  
Chemistry and  
Biochemistry.  
made to includ  
the biochemical  
lete ones, excep

The terminol  
ber of character  
tion of entries.  
of terms from  
try, by its very  
sciences. For th  
sciences as ch  
virology, biop  
been included i  
acteristic is the  
both standard  
are included to  
literature and t  
some of the no  
come standard  
characteristic i  
mous expression  
other only by  
onymous natur  
may not always  
cipal synonymo

ites a second mole-  
gy transfer, and the  
es the excitation en-

ER. 2. ALLERGEN.

initial and harmless  
to an animal which,  
injection, will trigger

avidson model.

-Davidson model.

ibody-like receptor  
ucing cell by which  
stimulate the cell to

obtained from whole  
with carbon dioxide

is The hypothesis  
systems I and II of  
parate systems such  
ossible only between  
system, but not be-  
one system to those  
llover hypothesis.

tical ultracentrifuge  
eparation and recove-  
e component having  
n coefficient.

io of the retention  
e retention volumes,  
are separated by gas

electrophoresis.

substances that are  
s, to carrier amphoteric  
isoelectric focusing  
ncing a relatively flat  
ent and thereby im-  
f components.

a group of cross-  
in gel filtration.  
a group of agarose

thogenic microorgan-  
the blood or in the

ide that has a seven-

f a septanose.

meable cell junction  
junction but occurs  
differs from a tight  
unctional proteins are  
ular fashion and that  
actually bringing the  
into direct contact.

o, sepsis.

**septicemia** The presence of pathogenic micro-organisms in the blood.

**septum** (*pl* septa; septums) A wall or a membrane that divides a cavity.

**Sequenase** Trademark for an enzyme preparation used in DNA sequencing. The enzyme is derived from bacteriophage T7 DNA polymerase and has been modified to improve its properties for sequencing.

**sequenator** An instrument for the automatic determination of amino acid sequences in a polypeptide chain; operation of the instrument is based on the repetitive application of the Edman degradation. *Aka* sequencer.

**sequence** 1. The linear order in which monomers occur in a polymer; the order of amino acids in a polypeptide chain, and the order of nucleotides in a polynucleotide strand are examples. 2. METABOLIC PATHWAY.

**sequence complexity** See complexity.

**sequence gap** A segment, consisting of one or more amino acids, that appears to be missing from a polypeptide chain when this chain is compared with others of the same protein but isolated from different sources, and when the chains are matched up so as to provide a maximum of sequence homology.

**sequence homology** The identity in sequence of either the amino acids in segments of two or more proteins, or the nucleotides in segments of two or more nucleic acids.

**sequence hypothesis** The hypothesis that the sequence of nucleotides in a nucleic acid specifies the sequence of amino acids in a protein.

**sequence isomer** One of two or more polymeric isomers that differ from each other in the sequence of the monomers in the chain.

**sequence polymer** A synthetic polypeptide consisting of identical repeating units, each of which is composed of more than one type of amino acid; the polymer (gly-ala-ser-val)<sub>n</sub> is an example. *See also* polyamino acid.

**sequencer** SEQUENATOR.

**sequence rules** RS SYSTEM.

**sequence specificity** The selectivity of a nuclease that accounts for its reaction being limited to specific base sequence in the nucleic acid.

**sequencing** The determination of the order of amino acids in a peptide, polypeptide chain, or protein, or the determination of the order of bases (nucleotides) in a nucleotide, polynucleotide strand, or nucleic acid.

**sequencing gel** A long, thin polyacrylamide gel slab used for nucleic acid sequencing.

**sequential feedback inhibition** The inhibition that is produced when one or more end products inhibit an enzyme in a metabolic pathway and the metabolite that accumulates as a

result of this inhibition then inhibits the first enzyme in the sequence and thereby shuts off the entire pathway.

**sequential induction** Enzyme induction in which a single inducer brings about the synthesis of a number of inducible enzymes; the first enzyme induced acts on the inducer, thereby transforming it into an inducer for the second enzyme, which in turn acts on the second inducer, and so on. *See also* coordinate induction.

**sequential mechanism** The mechanism of an enzymatic reaction in which two or more substrates participate in such a fashion that all the substrates must become bound to the enzyme before any products can be released. The mechanism is ordered if the substrates add to, and the products leave, the enzyme in an obligatory sequence; the mechanism is random if the substrates add to, and the products leave, the enzyme in a nonobligatory sequence.

**sequential model** A model for allosteric enzymes, proposed by Koshland, Nemethy, and Filmer, according to which the enzyme undergoes a series of conformational changes as the various ligands become bound to the enzyme. Different types of site interactions may occur of which symmetry preservation, as in the concerted model, may be a special case. In general, however, the symmetry of the enzyme molecule is not conserved, since a subunit changes its conformation as a ligand becomes bound to it. The capacity of the enzyme to bind substrate, positive effectors, and negative effectors is altered by the conformational changes which the subunits undergo. *Abbr* KNF model.

**sequential reactions** CONSECUTIVE REACTIONS.

**sequester** To form a chelate.

**sequestering agent** CHELATING AGENT.

**sequestration** CHELATION.

**sequestrene** ETHYLENEDINITROLOTETRAACETIC ACID.

**sequon** An obligatory sequence of amino acids that is required for a specific reaction. The term has been used to describe the tripeptide asn-x-thr or asn-x-ser that must occur in a protein for the asparagine (asn) to be able to act as a site of attachment for a carbohydrate moiety, thereby giving rise to a glycoprotein. *Var sp* sequeon.

**Ser** 1. Serine. 2. Seryl.

**SER** Smooth endoplasmic reticulum.

**serendipity** The gift for discovering valuable or useful things not specifically sought but recognized in the process of dealing with other things.

**serial dilution** The systematic and progressive dilution that is frequently used in immuno-